In-plant Study Demonstrating the Efficacy of Multiple Interventions to Demonstrate Overall Reduction of Microbial Load

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INTRODUCTION

Surrogate strains of *E.coli* that are non-pathogenic and mimic the behavior of pathogenic *E.coli* were developed by researchers to test the effectiveness of in-plant treatments in beef production facilities (1). While treatments can be tested for their efficacy within a laboratory setting, testing within the actual processing facility gives a much more accurate representation of overall cleanliness (2). Every possible condition within a processing facility cannot be accurately replicated within a laboratory, so it is critical to verify treatment effectiveness within the processing facility that will be utilizing the intervention. Currently, there exists no industry standard for the measuring of the 5-log reduction in raw products, and this project looked to validate the treatments applied to beef carcasses that reduce microbial load and create that standard operational procedure for such testing and validation.

OBJECTIVE

This in-plant study was conducted to create a standard operational procedure for the validation of antimicrobial treatments within a beef processing facilities using surrogate strains of bacteria. This study also compared the level of microbial attachment at different inoculation sites, as well as comparing two mediums used to swab carcasses.

MATERIAL AND METHODS

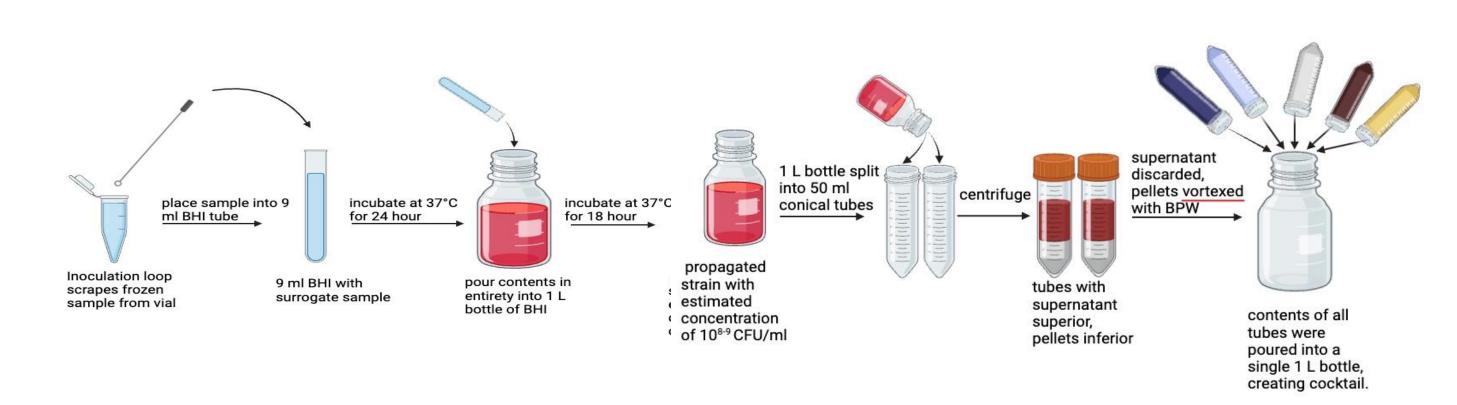


Figure 1: : Initial propagation of surrogate strains of E. coli strains BAA 1427-1431. Strains were propagated from frozen cultures, grown in BHI bottles, centrifuged, and combined to create a cocktail that was frozen at the Texas Tech University

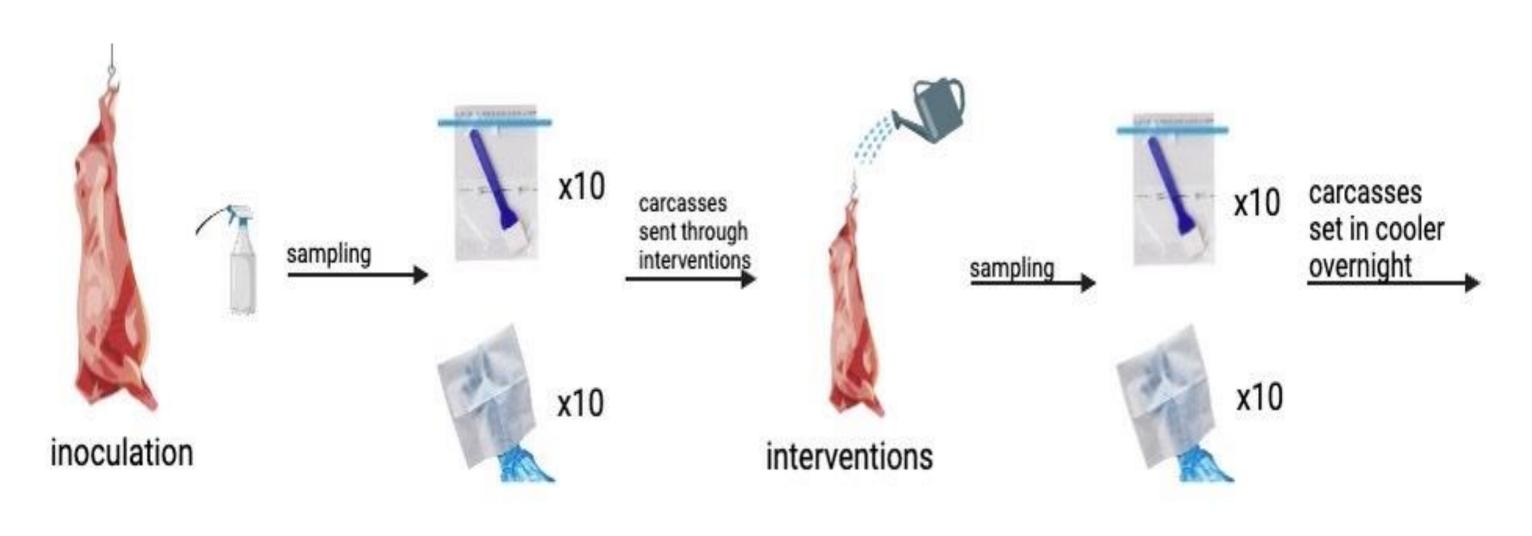
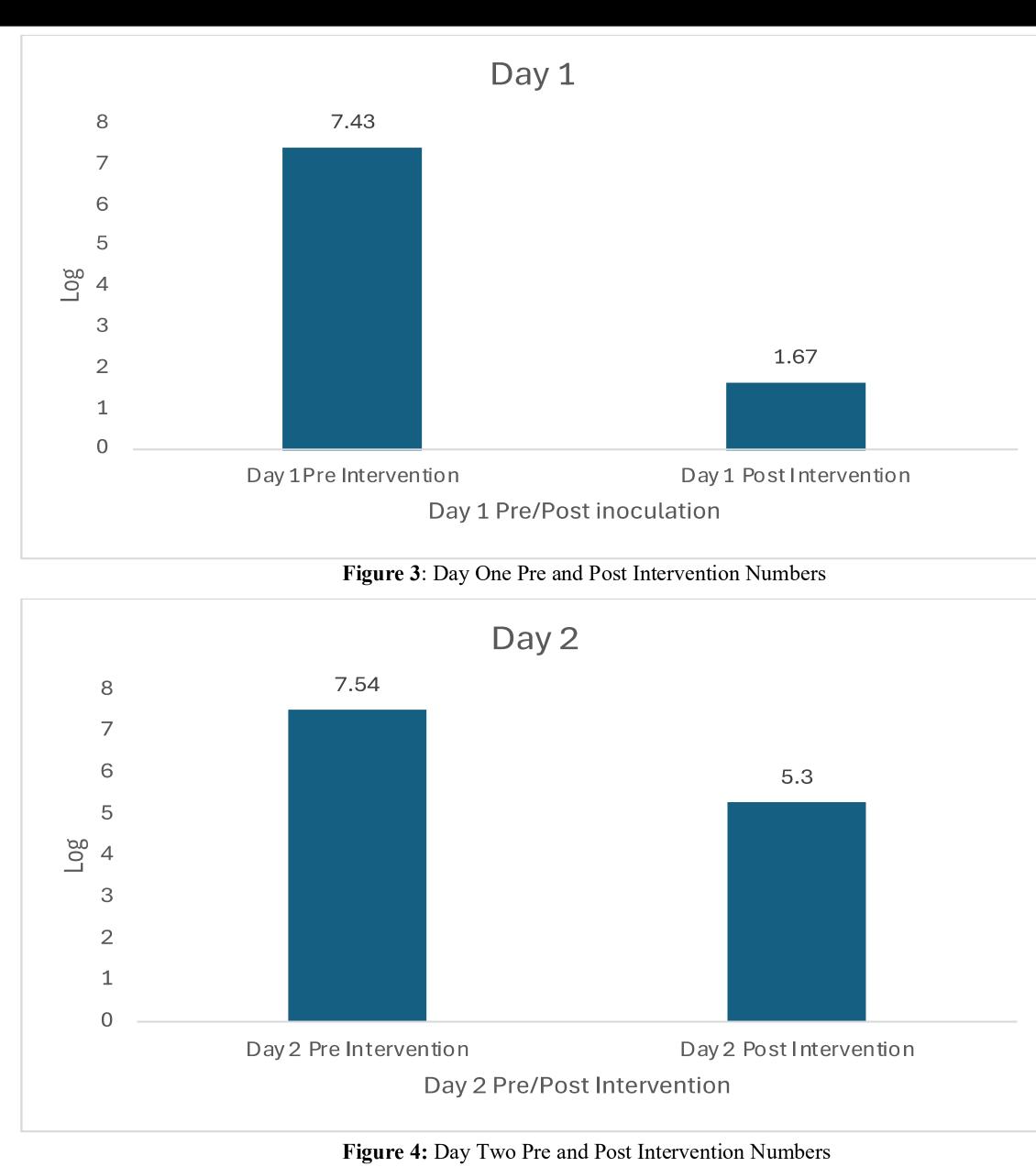


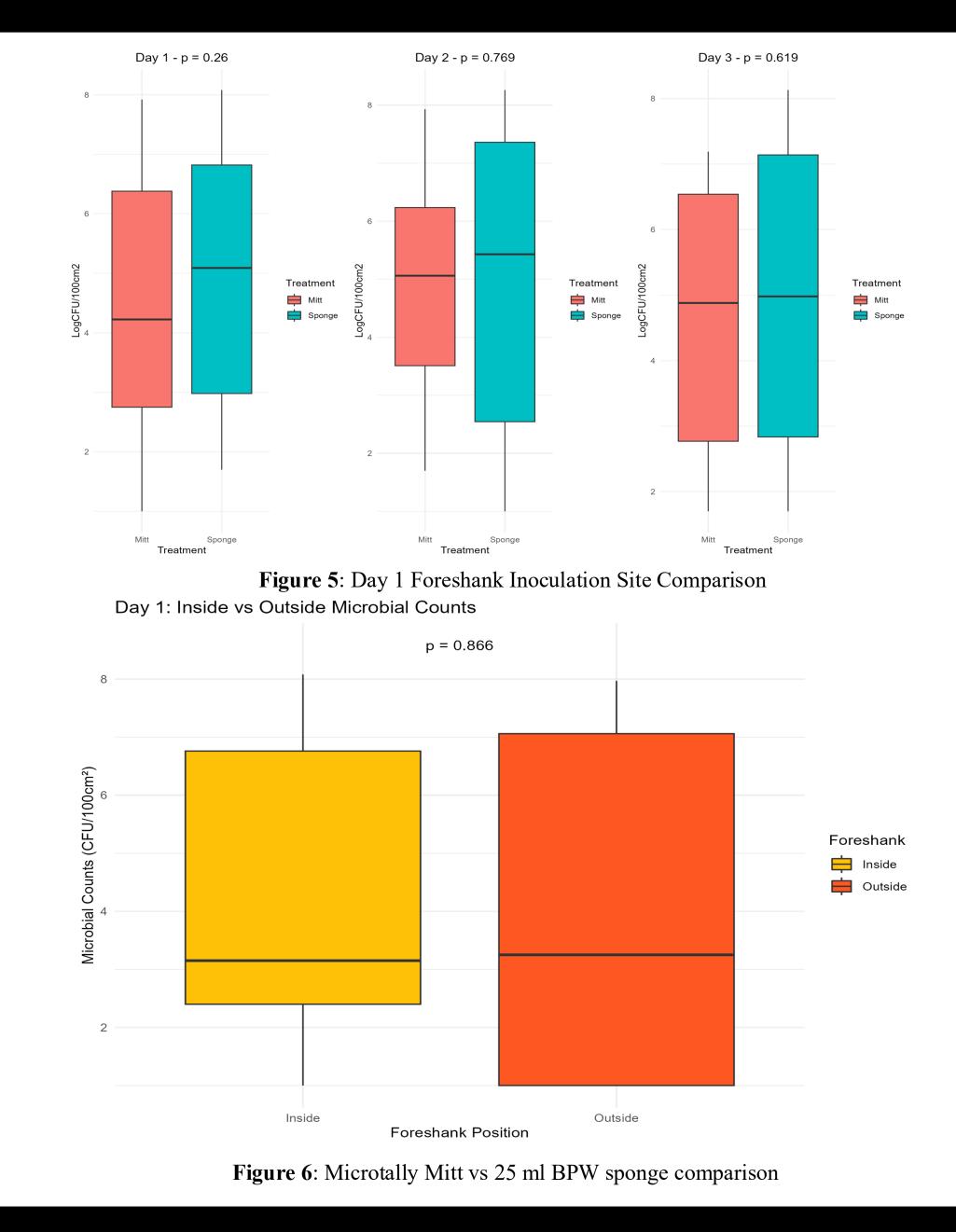
Figure 2: Day 1 Sampling Procedure: Day 1 sampling procedure followed to successfully inoculate and collect samples from foreshanks containing the applied surrogate cocktail.

RESULTS

attachment levels were with interventions yielding a 5.76 log reduction. attachment Day levels averaged to 7.54 log, interventions yielding a 2.24 log reduction, resulting in a 8.0 total achieved reduction when of combining both days treatments. When comparing a MicroTally mitt against a 25 ml prehydrated sponge, no statistical difference was (p=0.255).observed When comparing inoculation sites, the foreshank showed inner difference, statistical combined day one and day two data yielded a p-value of 0.677.







CONCLUSIONS

Based on the findings in this study, an 8-log reduction from all combined interventions was achieved to prevent microbial growth and kill potential contaminants. The sampling showed a consistent level of microbial reduction, with day one treatment clearly showing a higher level of effectiveness than day two. Sponge vs MicroTally cloth comparisons showed no statistical difference, as did the inner and outer foreshank comparision. All in all, the operational procedure replicated during each repetition yielded consistent results for a clear demonstration of intervention efficacy. This standard of operation is replicable industry wide for antimicrobial treatment validation.

REFERENCES

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